



# California Drug Recall Information



## Recall Name

**Qualitest Expands Recall of Hydrocodone Bitartrate and Acetaminophen Tablets, USP 10mg/500mg, Due to the Potential for Oversized Tablets**

Recall Date	Product Description	Recalling Firm	Recall Reason
12/06/12	Hydrocodone Bitartrate and Acetaminophen Tablets, USP 10mg/500mg  NDC# 0603-3888-multiple  <a href="#">List of affected NDC #s</a>	<b>Qualitest Pharmaceuticals, Inc.</b> Huntsville, AL	<i>Potential for Oversized Tablets</i>
Recall Class	Product Identification	Distribution	Affected Dates
I	Hydrocodone Bitartrate and Acetaminophen Tablets, USP 10mg/500mg, 100 count  Suspect Lots Recalled: <ul style="list-style-type: none"><li>Lot numbers beginning with the letter “C”</li></ul> Tablets are pink, capsule-shaped tablets with “ <b>3600</b> ” debossed on one side of the tablet and “ <b>V</b> ” on the other.	<b>CA</b> , nationwide	Distributed between February 20, 2012 and November 19, 2012

FOR ADDITIONAL INFORMATION, PLEASE VISIT:

<http://www.fda.gov/Safety/Recalls/ucm331218.htm>